

DETERGENT COMPOSITIONS FOR CLEANING

This invention relates to cleaning formulations based on water and water-immiscible organic substances which contain acrylic acids copolymers and vinyl esters as emulsifying agents.

BACKGROUND

5 Formulations designed for cleaning various surfaces (such as furniture and floors) and clothes contain numerous components with different chemico-physical properties. These formulations often contain raw materials which are miscible with one another with difficulty, or tend to separate.

10 It is common practice to use surfactants to homogenise multiphase systems (water-in-oil or oil-in-water emulsions) or to solubilise substances which are not miscible with one another.

15 Non-ionic surfactants, anionic, cationic or amphoteric surfactants or mixtures thereof are used for this purpose. Examples of non-ionic surfactants are alkylphenols, preferably octyl or nonylphenols ethoxylated with 1 to 25 moles of ethylene oxide, C₆-C₂₂ linear- or branched fatty alcohols ethoxylated, propoxylated or etho-propoxylated with 1 to 40 moles of ethylene oxide, propylene oxide or mixtures thereof, C₆-C₂₂ linear- or branched fatty acids and their amides, ethoxylated with 1 to 40 moles of ethylene oxide, propylene oxide or mixtures thereof, and C₆-C₂₂ linear- or branched fatty amines 20 ethoxylated, propoxylated or etho-propoxylated with 1 to 40 moles of ethylene oxide, propylene oxide or mixtures thereof.

Non-ionic surfactants with an HLB (Hydrophile-Lipophile Balance) value lower than 8 are classified as water-in-oil emulsifiers. Surfactants with an HLB value of between 8 and 14 are classified as detergents and wetting 25 agents. Surfactants with an HLB<14 are classified as oil-in-water emulsifiers and solubilising agents.

Examples of anionic surfactants are alkyl or alkenyl sulphates with the formula $R-O-SO_3^-M^+$, wherein R may have 10 to 22 carbon atoms and M is a cation of alkaline metal, ammonium or alkanolammonium, alkyl or alkenyl sulphates with the formula $R-O-(CH_2-CH_2)_n-SO_3^-M^+$, wherein R may have 10 to 22 carbon atoms, M is a cation of alkaline metal, ammonium or alkanolammonium, and n may have a value between 1 and 25, ethoxylated and sulphated alkylphenols, alkylpolysaccharide and alkylpolyglucoside sulphates, alkylbenzene sulphonates, primary or secondary C_6-C_{24} alkane sulphonates, C_6-C_{24} olefin sulphonates, sulphosuccinates and alkylsulphosuccinates, amides 5 of fatty acids with taurine and methyl-taurine wherein the sulphonic groups are generally salified with cations of alkaline metals, ammonium or alkanolammonium, soaps, with the carboxyl group in either the primary or the secondary position, alkyl ethoxy or (poly)ethoxy carboxylates with the general formula $RO(CH_2CH_2O)_xCH_2COOM$, wherein M is the cation of alkaline 10 metals, ammonium or alkanolammonium, sarcosinates with the general formula $R-CON(R_1)CH_2-COOM$, wherein M is the cation of alkaline metals, ammonium or alkanolammonium, and amphoteric surfactants. 15

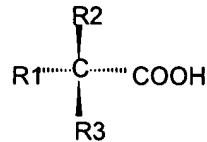
These surfactants are generally used in quantities of not less than 0.3% (w/w) and not more than 5% (w/w) of the weight of the total formulation when 20 used as a stabiliser, and in larger quantities when they are also used as a detergent, such as in washing machine detergent formulations.

DESCRIPTION OF THE INVENTION

It has now been found that copolymers of carboxylic acids containing an olefinic bond, with vinyl esters of aliphatic acids can wholly or partly 25 replace the conventional surfactants mentioned above, with the advantage that the formulation is more stable over time.

The polymers which can be used according to the invention are preferably copolymers of acrylic or methacrylic acid with vinyl esters of

branched aliphatic acids having the formula:



wherein R₁, R₂ and R₃, which are equal or different, represent linear or branched alkyl groups, with the proviso that the sum of the total carbon atoms, including carboxyl, is between 5 and 10, said copolymers being optionally 5 cross-linked with allyl ethers of polyvalent alcohols such as pentaerythritol triallyl ether.

The polymers described above are known, for example, from IT 1,200,111 and EP 724,875, and are used in the form of their alkaline metal, ammonium or amine salts.

10 These polymers mainly perform their activity in cases in which formulations based on water and water-immiscible organic solvents need to be stabilised, and also in cases where mixtures of polar organic solvents such as methanol, ethanol and dimethylformamide need to be stabilised with other solvents immiscible with them.

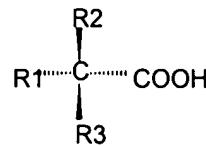
15 A characteristic of these copolymers is that not only they give the formulation containing them high viscosity, but also maintain said viscosity in the presence of high quantities of electrolytes such as sodium chloride, sodium sulphate, ammonium sulphate, ammonium phosphate, colorants and anionic surfactants. For this reason these copolymers are used in all products containing 20 a high electrolyte concentration.

DETAILED DESCRIPTION OF THE INVENTION

For the purposes of the present invention, the preferred copolymer consists of:

25 a) 98% to 80% of one or more carboxylic acids containing a double olefinic bond;

b) 2% to 20% of one or more vinyl esters of trialkylacetic acids having the formula:



wherein R₁, R₂ and R₃, which are equal or different, represent linear or branched alkyl groups, with the proviso that the sum of the total carbon atoms,

5 including the carboxyl, is between 5 and 10;

c) optionally between 0% and 3% of one or more cross-linking agents constituted by allyl ethers of polyvalent alcohols, calculated on the sum of the monomers referred to in paragraphs a) and b).

The copolymer in which component a) is acrylic acid, component b) is 10 isodecanoic acid vinyl ester and component c) is pentaerythritol triallyl ether is particularly preferred for the purposes of this invention.

According to the invention, the quantity of copolymer used is between 0.01% and 10%, and preferably between 0.1% and 3%, of the total weight of the formulation.

15 Formulations obtained by using a copolymer according to the invention as stabiliser and/or emulsifier, alone or mixed with surfactants which are commonly employed according to the state of the art, are particularly stable, and may take the form of water-in-oil or oil-in-water emulsions, or a gel.

Examples of said formulations are emulsions containing pine oil, 20 limonene, silicones, liquid paraffin, waxes, designed for cleaning furniture or floor or clothes. Moreover, the copolymers to which this invention relates are particularly suitable to stabilise formulations containing silicone-based anti-foaming agents or fragrances, constituents which are difficult to solubilise and have a strong tendency to separate. Anti-foaming agents are present in the 25 formulations according to the invention in percentages of between 0.01 and

40% of the total weight of the formulation.

When present, essential oils or fragrances are used in concentrations of between 0.01 and 50% of the total weight of the formulation.

The quantity of copolymer used in said formulations varies according to 5 the nature and quantity of the elements to be emulsified and it is comprised between 0.01% and 10%.

Before use, the copolymer must be pre-dispersed in one or more of the 10 phases present in the formulation, usually the aqueous phase or the oily phase; after mechanical homogenisation of the phases, the pH is adjusted to the 15 desired value with an alkaline substance such as sodium hydroxide, potassium hydroxide, an alkanolamine, an aliphatic amine, aminoethylpropanol, aminoethyl propanediol, etc..

The examples set out below illustrate some compositions of 15 formulations containing the copolymer and/or a mixture of copolymer and surfactant in a smaller quantity than traditional formulations.

In the examples, the copolymer according to the invention is indicated as COPOLYMER. This copolymer was prepared according to example no. 2/F of IT 1,200,111.

EXAMPLE 1

20 Compositions 1, 2 and 3 describe a floor wax in the form of an emulsion containing both synthetic wax and conventional natural wax.

This formulation was prepared with surfactant acting as emulsifier alone (Formula 1), with a mixture of surfactant acting as emulsifier and COPOLYMER (Formula 2), or with the COPOLYMER alone, wholly 25 replacing the surfactant acting as emulsifier (Formula 3).

In the product with surfactant only, separation begins to occur after 2 months at 40°C, whereas the two formulations containing the COPOLYMER show excellent stability over time and increased viscosity. All the viscosities

were measured at 20°C with a Brookfield RV viscosimeter, spindle 2.

Composition 1 - Flo. r wax (comparison formula)	
Carnauba wax (natural wax)	2%
Sasolwax A2 (synthetic wax)	2%
Emcol 4100M (surfactant based on disodium myristamide MEA sulphosuccinate, acting as emulsifier)	1.5%
Empilan KCL 11 (surfactant based on ethoxylated fatty alcohol 11EO)	0.3%
Perfume	0.3%
Germaben II (preservative)	0.2%
NaOH (10%)	at pH 7
Demineralised water	q.s. for 100%

pH: 7

Viscosity (mPa.s): 80

Composition 2 – Floor wax (according to the invention)	
Carnauba wax (natural wax)	2%
Sasolwax A2 (synthetic wax)	2%
Emcol 4100M (surfactant based on disodium myristamide MEA sulphosuccinate, acting as emulsifier)	0.5%
Empilan KCL 11 (surfactant based on ethoxylated fatty alcohol 11EO)	0.3%
COPOLYMER	0.5%
Perfume	0.3%
Germaben II (preservative)	0.2%
NaOH (10%)	at pH 7
Demineralised water	q.s. for 100%

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pH: 7

Viscosity (mPa.s): 700

Composition 3 – Floor wax (according to the invention)	
Carnauba wax (natural wax)	2%
Sasolwax A2 (synthetic wax)	2%
COPOLYMER	1%
Empilan KCL 11 (ethoxylated fatty alcohol 11EO)	0.3%
Perfume	0.3%
Germaben II (preservative)	0.2%
NaOH (10%)	at pH 7
Demineralised water	q.s. for 100%

pH: 7

Viscosity (mPa.s): 700

	Composition 1	Composition 2	Composition 3
Initial viscosity (mPa.s)	80	700	800
Viscosity after 1 month at 40°C (mPa.s)	80	700	800
Viscosity after 2 months at 40°C (mPa.s)	70 (slight separation)	750	870
Viscosity after 3 months at 40°C (mPa.s)	70 (separation into two phases)	750	900

EXAMPLE 2

Compositions 4, 5 and 6 below describe a cream for hard surfaces containing a large quantity of d-Limonene. This formulation is traditionally prepared using a surfactant acting as emulsifier (Composition 4). In formula 5, part of the surfactant acting as emulsifier is replaced by the copolymer, while Composition 6 only contains the copolymer as emulsifying agent.

Composition 4 - Cream with d-Limonene (comparison formula)	
d-Limonene	8%
Amphisol K (surfactant based on potassium cetyl phosphate, acting as emulsifier)	1.55%
Lutensol (surfactant based on ethoxylated fatty alcohol 7EO)	2%
Nansa (dodecyl benzenesulphonic acid)	0.5%
Propylene glycol	3.5%
Caustic soda	at pH 7.5
Demineralised water	q.s. for 100

pH: 7.5

Viscosity (mPa.s): 100

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Composition 5 - Cream with d-Limonene (according to the invention)	
d-Limonene	8%
COPOLYMER	0.55%
Amphisol K (surfactant based on potassium cetyl phosphate, acting as emulsifier)	0.5%
Lutensol AO7 (surfactant based on ethoxylated fatty alcohol 7EO)	2%
Nansa (dodecyl benzenesulphonic acid)	0.5%
Propylene glycol	3.5%
Caustic soda	at pH 7.5
Demineralised water	q.s. for 100

pH: 7.5

Viscosity (mPa.s): 1000

Composition 6 - Cream with d-Limonene (according to the invention)	
d-Limonene	8%
COPOLYMER	0.55%
Lutensol AO7 (surfactant based on ethoxylated fatty alcohol 7EO)	2%
Nansa (dodecyl benzenesulphonic acid)	0.5%
Propylene glycol	3.5%
Caustic soda	at pH 7.5
Demineralised water	q.s. for 100

pH: 7.5

Viscosity (mPa.s): 1000

	Composition 4	Composition 5	Composition 6
Initial viscosity (mPa.s)	100	1000	1000
Viscosity after 1 month at 40°C (mPa.s)	100	1100	1100
Viscosity after 2 months at 40°C (mPa.s)	150	950	1050
Viscosity after 3 months at 40°C (mPa.s)	170 (separation into two phases)	925	1050

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EXAMPLE 3

Compositions 7 and 8 are two washing machine detergent formulations containing a silicone-based anti-foaming agent. Composition 8 shows excellent stability over time, whereas Composition 7 leads to complete separation of the silicone from the rest of the formulation.

Composition 7 – HEAVY-DUTY LIQUID (comparison example)	
Nansa (dodecyl benzenesulphonic acid)	8%
Empimin KSN 70 LA (lauryl ether sulphate 3EO)	7%
Lutensol AO7 (surfactant based on ethoxylated fatty alcohol 7EO)	3%
Edenor K12-18 (mixture of fatty acids of tallow/coconut)	2%
KOH (50%)	1.2%
Bidistilled olein	1%
Anti-foaming agent (silicone-based)	0.4%
Citric acid monohydrate	2%
EDTA (sequestering agent)	0.06%
NaOH (10%)	at pH 9
Demineralised water	q.s. for 100%

pH: 9

Viscosity (mPa.s): 200

Composition 8 – HEAVY-DUTY LIQUIDS (according to the invention)	
Nansa (dodecyl benzenesulphonic acid)	8%
Empimin KSN 70 LA (lauryl ether sulphate 3EO)	7%
Lutensol AO7 (surfactant based on ethoxylated fatty alcohol 7EO)	3%
Edenor K12-18 (mixture of fatty acids of tallow/coconut)	2%
COPOLYMER	0.5%
KOH (50%)	1.2%
Bidistilled olein	1%
Anti-foaming agent (silicone-based)	0.4%
Citric acid monohydrate	2%
EDTA (sequestering agent)	0.06%
NaOH (10%)	at pH 9
Demineralised water	q.s. for 100%

	Composition 7	Composition 8
Initial viscosity (mPa.s)	200	1000
Viscosity after 1 month at 40°C (mPa.s)	220	1100
Viscosity after 2 months at 40°C (mPa.s)	180 (separates)	950
Viscosity after 3 months at 40°C (mPa.s)	180 (separates)	950